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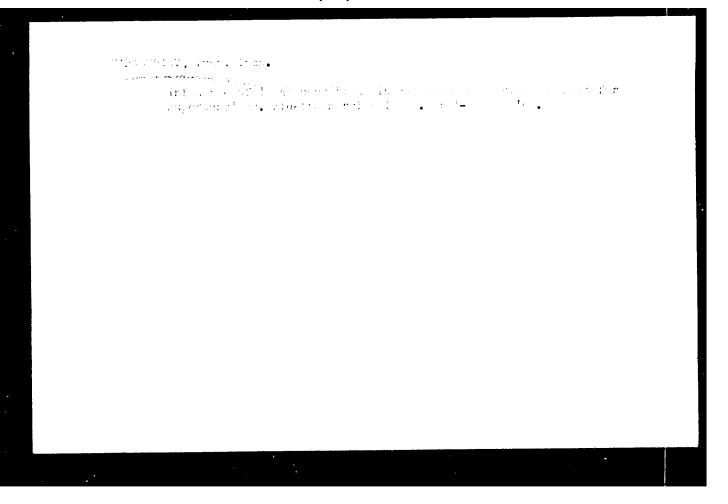
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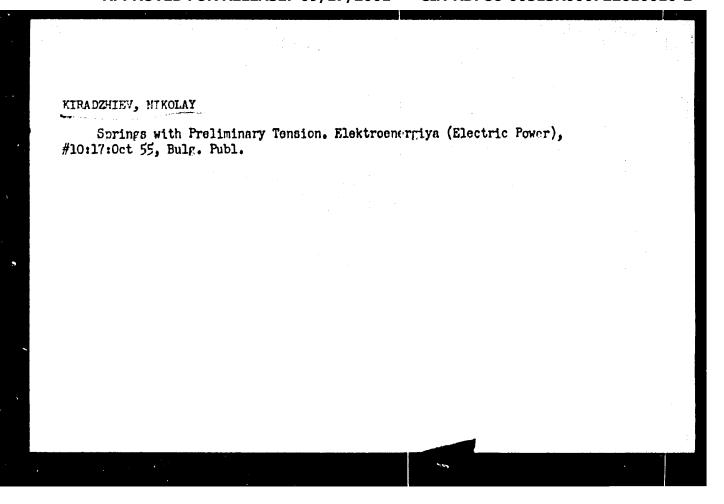


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Dissertation for Candidate of Technical Sciences, All-Union Sci. Res. Inst. of Water-Supply, Sewerage, Hydraulic-Engineering Structures and Engineering Hydrogeology (VODGEO)

Subject: Hydroengineering building and construction

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SO: Sum. No. 480, 9 May 55

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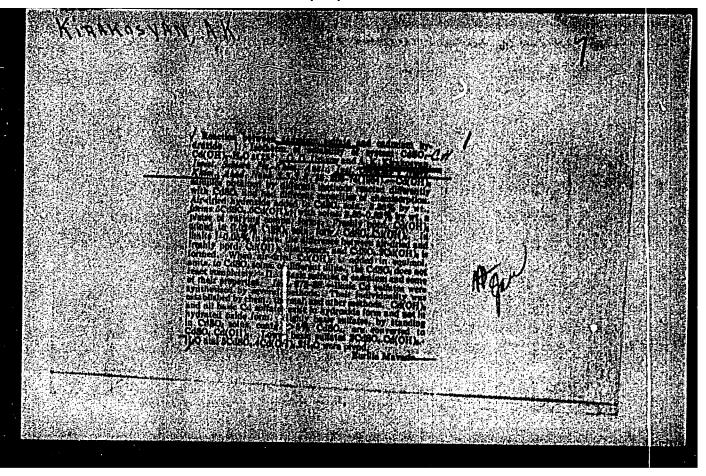
Preparation of cadmium-sulfate ammines and cadmium-ammonium sulfate from aqueous solutions. Report no.2: Solubility isothern of the ternary system CdSO<sub>4</sub> • (NH<sub>4</sub>)2SO<sub>4</sub> — NH<sub>3</sub> — H<sub>2</sub>O at 25°. Izv.Sekt.fiz.-khim.anal. 22:261-271 53. (MLRA 7:5)

1. Institut obshchey i neorganicheskoy khimii im. N.S.Kurnakova Akademii nauk SSSR. (Ammines) (Sulfates) (Systems (Chemistry))

Production of cadmium sulfats ammine and cadmium ammonium sulfate from aqueous solutions. III. Solubility isotherny of the system C4, NH4804, OH-Ho at 25°. Co. Urraya and A. Kitalovyan, Irsul-Seletora Fis.-Rhim. And., Akdd. Nanh S.-S.-R. 23, 249-57 (1053).—In the systems (NII), SO<sub>4</sub>-NII-H<sub>2</sub>O and Cd. (011),-NH<sub>2</sub>-H<sub>2</sub>O no chem. compris. were formed. In the system ClSO<sub>4</sub>-(NII), SO<sub>4</sub>-H<sub>2</sub>O the sol. CdSO, (NII), SO<sub>4</sub>-Olifor in the fisher pind. Cd(OH), was formed. In the system ClSO<sub>4</sub>-Cd(OH),-H<sub>2</sub>O, when freshly ppid. Cd(OH), was used, CdSO<sub>4</sub>-Cd(OH),-H<sub>2</sub>O, CdSO<sub>4</sub>-2Cd(OH), were formed. The latter formed when the CdSO<sub>4</sub>-2NH<sub>4</sub>OH is divided into 2 parts. A metathetical reaction first takes place. As the NH4, concn. increases [Cd, (NII<sub>4</sub>)(H<sub>2</sub>O);SO<sub>4</sub> is formed up to 12.5% NHe. Purther increase in NH4, when the soln, is said, with CdSO<sub>4</sub>-causes saiting out of the complex. Study of other sections on the diagram revealed the formation of numerous complexes within the system. Along the diagonals of the system no stable complex. Along the diagonals of the system no stable complex ending the diagonals of the system no stable complex are formed. The system comprises 7 crystin. fields: CdSO<sub>4</sub>,8/3H<sub>2</sub>O, CdSO<sub>4</sub>,(NH4),SO<sub>6</sub>, Cd(NH4), Cd(OH), Mil<sub>2</sub>O<sub>5</sub>, (NH4),SO<sub>6</sub>, [Cd(NH4), Cd(OH), Mil<sub>2</sub>O<sub>6</sub>, (NH4),SO<sub>6</sub>, [Cd(NH4), Cd(OH), Mil<sub>2</sub>O<sub>6</sub>, (NH4),SO<sub>6</sub>, [Cd(NH4), Mil<sub>2</sub>O<sub>6</sub>, (NH4),SO<sub>6</sub>, [Cd(OH), Mil<sub>2</sub>O<sub>6</sub>, (NH4),SO<sub>6</sub>, [Cd(OH), Mil<sub>2</sub>O<sub>6</sub>, [Cd(OH), Mil<sub>2</sub>O<sub>6</sub>,



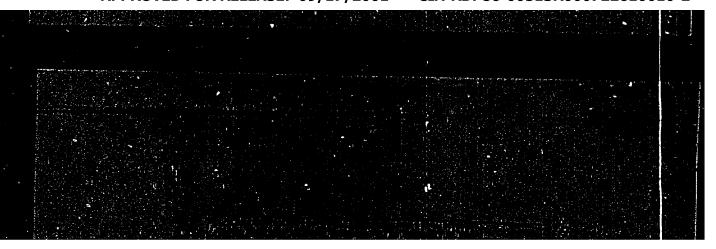
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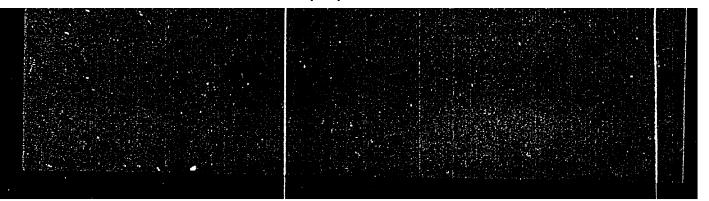
URAZOV, G.G.; KIRAKOSYAN, A.K.

Study of the reaction of cadmium sulfate with cadmium hydroxide. Report no.2. Basic cadmium sulfates and certain of their properties. Isv. Sekt.fiz.-khim.anal. no.25:275-288 154. (MIRA 8:5)

1. Institut obshchey i neorganicheskoy khimii im. N.S.Kurnakova Akademii nauk SSSR. (Cadmium sulfate)



"APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722610010-2



Kirkosulatry

Card 1/1

Pub. 22 - 31/54

Authors

Urasov, G. O., Acidemician, and Kirakosyan, A. K.

Title

Reaction between ammonia and divalent metal halides in an aqueous medium Solubility of cadmium chloride in aqueous ammonia solutions

Periodical

Dok. AN SSSR 106/2, 290-293, Jan 11, 1956

Abstract

The reaction between cadmium chloride and ammonia in an aqueous medium was investigated by the isothermal solubility method. The zone of crystallization of the hydroxide residues and the ammoniates was studied at a temperature range of from 0 to 25°. It was found that the solubility of cadmium chloride in the zone of crystallization of higher CdCl<sub>2</sub> ammoniates depends upon temperature. The changes in the composition of compounds crystallizing at 25° are explained. Nine references: 1 Russ. 5 Cerm. and 3 French (1842-1912). Diagrams.

Institution :

Acad. of Sc., USSR, Inst. of Gen. and Inorgan. Chem. im. N. S. Kurnakov

Submitted

September 30, 1955

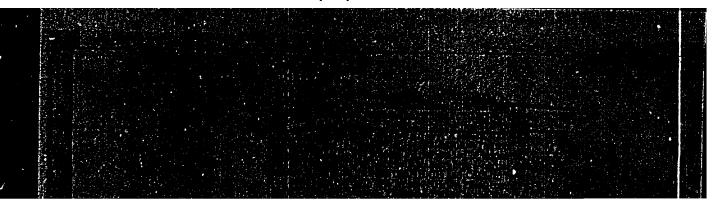
URAZOV, G.G. [decoased]: KIRAKQSYAN, A.K. GALUSTYAN, V.D.

FIRMKHOSAN A C

Interaction between armonium and the halides of divalent metals in an aqueous medium. Part 2: Solubility of cupric bromide in aqueous solutions of ammonia. Zhur. neorg. khim. 2 no.5:1105-1114 Mv 157. (MLRA 10:10)

1. Institut obshchey i neorganicheskoy khimii imeni N.S. Kurnakeva. Akademii nauk SSSR.

(Copper browides) (Ammonia)



20-114 3-31/60

AUTHORS:

Urazov, G. G., Member of the AN USSR, Kirakosyan, A. K.,

Mkhitaryan, R. S.

TITLE:

An Investigation of the Interaction Between Ammonia and Zinc Chloride in Water Solutions (Izucheniye vzaimodeystviya rezhdu ammiakom i khloristym takinkom v vodnoy srede)

PERIODICAL:

Doklady Akademii Nauk SSSR,1957, Vol 114, Nr 3, pp 564-567 (USSR)

ABSTRACT:

An investigation of the interaction between anmonia and zinc chloride in water solutions during an entire phase of the concentration of the solution of the latter at different temperatures has never been carried out. Anhydrous hexammine zinc chloride has already been obtained earlier by blowing gaseous ammonia through anhydrous zinc chloride. The expansion capacity of the dissociation of hexammine zinc chloride and its decomposition products was determined as well. Monoaquo-pentammine zinc chloride was produced by cooling a saturated ammonia solution of zinc chloride. Tetrammine zinc chloride with different water content was obtained by cooling the ammonia solution of zinc chloride. The decomposition tempera-

Carapproved for releasemon/17/2001. Digial-RDP86-00513B0Q0722610010-2"

An Investigation of the Interaction Between Ammonia and Zinc Chloride in Water Solutions

with a content of crystallized water or anhydrous, was produced by different methods: dissolution of zinc oxide in chloral ammonia solutions, blowing of ammonia gas through zinc chloride solution, thermal decomposition of higher ammonia compounds of zinc chloride, etc. The dissociation temperatures and the beginning of decomposition were determined. Monoammine zinc chloride is the final product resulting from thermal decomposition of higher ammonia compounds and can be distilled without disintegrating. There then follows an experimental part with description of methods of production. Discussion of results: The crystallization of basic salts containing ammonia is completed with a content of 9.04 % weight of ammonia and 18.92 % weight of zinc chloride in a fluid phase of equal weight. The zinc chloride content rises with increasing concentration of ammonia and is directly proportional to the content of the latter in the fluid phase. The crystallization of the basic ammonia salts of zinc chloride is the result of a partial rearrangement-reaction process between ammonia hydroxide and zinc chloride. With an increasing concentration of ammonia in the fluid phase the exchange between the two substances decreases according to

Card 2/4

An Investigation of the Interaction Between Ammonia and Zinc Chloride in

parabolic dependence. The solid phase of the crystallisation of basic salts of zinc chloride are structures of varying composition. They all, without exception, contain ammonia. Their composition is ZnCl<sub>2</sub>.nZn(OH)<sub>2</sub>.sNH<sub>3</sub>.xH<sub>2</sub>O. The coefficients

n, s and x are of different values, integer figures as well as fractions. The content of ammonia on the solid phase depends on its concentration in the fluid phase. The more of it contained in the fluid phase, the greater is its portion in the solid phase. The content of zino hydroxide decreases accordingly. In spite of different chemical composition, these salts possess the same properties (thermal, crystalloptical, etc.) and similar Debaille diagrams. Agreement of some properties as well as of the crystal lattice with their isomorphous group substitution, is probable as well of OH and NH, possibly also of H<sub>2</sub>O and NH<sub>2</sub>. Furthermore, the crystallization and solubility as well as the temperature curve (fusion and destillation) of the substances treated is described. There are 4 figures and 11 references, 1 of which is Soviet.

Card 3/4

AUTHORS: Urazov, G. G. (Deceased), Kirakosyan, A. K., 15thitaryan,

Investigations Concerning the Interaction Between Ammonia and Zine Salts in an Aqueous Medium (Izucheniye vzaimodeystviya mezhdu ammiakom i solyami tsinka v vodnoy srede)

I. The Solubility of Zine Chloride in Aqueous Ammonia Solutions (I. Rastvorimost khloristogo tsinka v vodnoammiachnykh rastvorakh)

PERIODICAL: Zhurnal Neorganicheskoy Khimii, 1958, Vol.3, Nr 2, pp.464-474 (USSR)

The solubility of zinc chloride in aquecus ammonia solutions of all concentrations of ammonia in the liquid phase was investigated. The solid phases of the system ZnCl<sub>2</sub>-NH<sub>2</sub>H<sub>2</sub>O at an ammonia-content of 9,04 % at 25 °C crystallize to basic salts of different composition. The composition of these basic salts can be expressed by the following general formula: ZnCl<sub>2</sub>·n Zn(OH)<sub>2</sub> · s NH<sub>2</sub> + x H<sub>2</sub>O, in which the coefficients n, s and x denote different values in integer or fractional

Investigations Concerning the Interaction Between Ammonia and Zinc Salts in Aqueous Medium. I. The Solubility of Zinc Chloride in Aqueous Anmonia

numbers. The basic salts were investigated thermally, crystallographically and by X-ray analysis. The crystal lization of the ammoniacal compounds in the system ZnCl<sub>2</sub>-NH<sub>2</sub>-H<sub>2</sub>O was performed at temperatures of O and 25°C. In the system ZnCl<sub>2</sub>-NH<sub>4</sub>-H<sub>2</sub>O at O°C two compounds of the following composition crystallize: ZnCl<sub>2</sub> · 2.2 NH<sub>2</sub> · 0.5 H<sub>2</sub>O and ZnCl<sub>2</sub> · 5.75 NH<sub>3</sub> · 0.75 H<sub>2</sub>O, and at 25°C: ZnCl<sub>2</sub> · 2.2 NH<sub>2</sub> · 0.5 H<sub>2</sub>O and ZnCl<sub>2</sub> · 5.35 NH<sub>3</sub> · 0.33 H<sub>2</sub>O. The crystallization of ZnCl<sub>2</sub> · 2.2 NH<sub>2</sub> · 0.5 H<sub>2</sub>O at O°C and 25°C begins at 10.96% and ends at 24.46% of ammonia in the aqueous phase. The crystallization of ZnCl<sub>2</sub> · 0.75 H<sub>2</sub>O at O°C and ZnCl<sub>2</sub> · 5.35 NH<sub>3</sub> · 0.33 H<sub>2</sub>O at 25°C begins at 24.64% ammonium in the aqueous phase. The crystallization proceeds irregularly and is dependent on the addition of ammonium in the aqueous phase. At a higher concentration the crystallization slows down. The thermographic analysis showed that in basic salts two endothermic effects occur, at 110-125°C dehydration occurs and at 220-225°C the hydroxide form is converted into oxide. The thermographic analyses of ZnCl<sub>2</sub>·2,2 NH<sub>2</sub>·0.5 H<sub>2</sub>O showed three endothermic effects: at 65°C and at 125°C -

Card 2/3

KirAKosyAn, A.K.

AUTHORS:

Urazov, C. C. (Deceased), Kirakosyan, A. K., 78-2-31/43 Mkhitaryan, R. S.

TITLE:

Investigations Concerning the Interaction Between Ammonia and Zinc Salts in an Aqueous Medium (Izucheniye vzaimodeystviya

mezhdu ammiakom i solyami tsinka v vodnoy srede)

II. The Solubility of Zinc Bromide in Aqueous Ammoniacal

Solutions (II. Rastvorimost' bromistogo tsinka v

vodnoammiachnykh rastvorakh)

PERIODICAL:

Zhurnal Neorganicheskoy Khimii, 1958, Vol. 3, Nr 2,

PP- 475-483 (USSR)

ABSTRACT:

The solubility of zinc bromide in aqueous ammonia solutions was determined. It is shown that up to 11,6% ammonia in the aqueous phase ammoniacal basic salts of zinc bromide of different composition crystallize. The interaction of ammonia and zinc bromide in an aqueous medium takes place in two phases: at a low concentration of ammonia, ammoniacal basic salts crystallize and at a higher concentration

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ammoniacates crystallize. The basic salts can be expressed by the following general formula:

Investigations Concerning the Interaction Between Ammonia 78-2-31/43 and Zinc Salts in an Aqueous Medium.

II. The Solubility of Zinc Bromide in Aqueous Ammoniacal Solutions

ZnBr<sub>2</sub> . n Zn(OH)<sub>2</sub> . s NH<sub>3</sub> . x H<sub>2</sub>O where the coefficients n, s and x may have different values (n varies between 0,9 - 3,2, s between 1,5 - 2,0). The thermographic analyses of the basic salts showed that ammonia escapes at 112, 135 and 240-250°C. A complete escape of ammonia by thermal analysis does not occur. The residue after treatment at 250°C has the following composition:

ZnBr<sub>2</sub> . 1,6 ZnO . NH<sub>3</sub>.

The X-ray analyses of the basic salts indicate a crystalline structure. At O°C ZnBr<sub>2</sub> . 4NH<sub>3</sub> and ZnBr<sub>2</sub> . 5,25 NH<sub>3</sub> . 0,5 H<sub>2</sub>O, crystallize from the system ZnBr<sub>2</sub>-NH<sub>3</sub>-H<sub>2</sub>O, and at 25°C - ZnBr<sub>2</sub> . 5,5 NH<sub>3</sub>. All ammoniacates are unstable compounds and decompose in air at room temperature. From the thermographic analysis of ZnBr<sub>2</sub> . 4NH<sub>3</sub> two endothermic effects are to be

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Investigations Concerning the Interaction Between Ammonia 78-2-31/43 and Zinc Salts in an Aqueous Medium.

II. The Solubility of Zinc Bromide in Aqueous Ammoniacal Solutions

seen: 1. at 145-155° C- corresponds to the escape of 1 mol. ammonium, the residue has a composition of ZnBr<sub>2</sub>. 3NH<sub>3</sub>; 2. at 240-250°C - again 1 mol. ammonium escapes and ZnBr<sub>2</sub>. 2NH<sub>3</sub> remains as residue. At a temperature higher than 250°C the decomposition of diamino-zinc-bromide occurs. The thermographic curves of ZnBr<sub>2</sub>. 25NH<sub>3</sub>. 0,5H<sub>2</sub>O and ZnBr<sub>2</sub>. 5,5NH<sub>3</sub> are equal. These curves have four endothermic effects. The first one occurs at 47-50°C under the escape of ammonium and the formation of ZnBr<sub>2</sub>. 5NH<sub>3</sub>, the second effect at 80°C under the giving off of 1,5% ammonium and the formation of ZnBr<sub>2</sub>. 4NH<sub>3</sub>. The two other effects are in agreement with the thermographic decomposition of tetramine-zinc-bromide. On heating of the ammoniacates, even at a temperature higher than 500°C, no complete escape of ammonium is attained.

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Investigations Concerning the Interaction Between Ammonia 78-2-31/43 and Zinc Salts in an Aqueous Medium.

II. The Solubility of Zinc Bromide in Aqueous Ammoniacal Solutions

There are 8 figures, 4 tables, and 6 references, 1 of which is Slavic.

SUBMITTED: February 7, 1957

AVAILABLE: Library of Congress

Card 4/4

KIRAKUSYAN.

AUTHORS:

Urazov, G. G., (Deceased), Kirakosyan, A. K., Mkhitaryan, P. S.

78-2-32/43

TITLE:

Investigations on the Interaction Between Zinc Salts and Ammonia in an Aqueous Medium (Izucheniye vznimodeystviya

mezhdu solyami tsinka i ammiakom v vodnov srede)

III. The Solubility of Zinc Iodide in Aqueous Ammoniacal Solutions (III. Rastvorimost' yodistogo tsinka v vodno-

ammiachnykh rastvorakh)

PERIODICAL:

Zhurnal Neorganicheskoy Khimii, 1958, Vol. 3, Kr 2, pp.484-

490 (USSR)

ABSTRACT:

The solubility of zinc iodide in aqueous ammoniacal

solutions at 0° and 25°C was determined. The crystallization

of basic salts in the system  $\text{ZnJ}_2\text{-NH}_3\text{-H}_2\text{O}$  terminates at

2,82% NH..

The ammoniacal basic salts of zinc iodide have the following general formula:  $ZnJ_2$ .  $nZn(OH)_2$ . s  $NH_3$ . x  $H_2O$ ,

where the coefficients n, s and x may have different values. At  $0^{\circ}$  in the system  ${\rm ZnJ_2-NH_3-H_20}$  the following ammoniacates

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Investigations on the Interaction Between Zinc Salts and 78-2-32/43 Ammonia in an Aqueous Medium III. The Solubility of Zinc Iodide in Aqueous Ammoniacal Solutions

orystallize:  $ZnJ_2$  · 4  $NH_3$  ·  $H_2O$  and  $ZnJ_2$  · 5  $NH_3$  ·  $H_2O$  , and at  $25^{\circ}C$  :  $ZnJ_2$  · 4  $NH_3$  ·  $H_2O$  and  $ZnJ_2$  · 5  $NH_3$  · 0,5  $H_2O$  ·

In the thermal analysis of the ammoniacal bisic salts dehydration occurs at loo and 135°C and at 205°C a conversion of zine hydroxide to zine exide and a partial escape of NH, is to be observed. In the thermographic analysis of the ammoniacates ZnJ. 5 NH<sub>3</sub>. H<sub>2</sub>O three endothermic effects occur: 1) at 35 = 50°C with giving off water; 2) at 100 - 105°C with giving off 1 mol. ammonium, ZnJ<sub>2</sub>.4NH<sub>3</sub>H<sub>2</sub>O remaining as a residue; 3) at 195 - 215°C under the formation of anhydrous triamine-zine-iodide ZnJ<sub>2</sub>. 3NH<sub>3</sub>.

In the decomposition of ZnJ<sub>2</sub>. 4 NH<sub>3</sub>. H<sub>2</sub>O two endothermic effects occur: 1) at 100 - 110°C with giving off coarse moisture; 2) at 195 - 215°C with giving off crystal water and 1 mol. ammonium and the formation of ZnJ<sub>2</sub>. 3NH<sub>3</sub>. On further heating the triamine decomposes.

The ammoniacates of zinc iodide are well crystallizable bodies and are difficult to dissolve in concentrated ammoniacal solutions. There are 6 figures, 3 tables, and

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#### CIA-RDP86-00513R000722610010-2 "APPROVED FOR RELEASE: 09/17/2001

Investigations on the Interaction Between Zinc Salts and Ammonia in an Aqueous Medium

78-2-32/43

III. The Solubility of Zinc Iodide in Aqueous Ammoniacal Solutions

7 references, 1 of which is Slavic.

SUPMITTED:

February 7, 1957

AVAILABLE:

Library of Congress

Card 3/3

WIRAKUSYAN AM

AUTHORS:

Urazov, G. G., (Deceased), Kirakosyan, A. K., Mkhitaryan, R. S.

78-2-33/43

TITLE:

Investigations Concerning the Interaction Between Zinc Salts and Ammonia in an Aqueous Hedium (Izucheniye vzaimodeystviya

mezhdu solyami tsinka i ammiakom v vodnov srede)

IV. The Solubility of Zinc Nitrate in Aqueous Ammoniacal Solutions (IV. Rastvorimost' azotnokislogo tsinka v

vodnoammiachnykh rastvorakh)

PERIODICAL:

Zhurnal Neorganicheskoy Khimii, 1958, Vol. 3, Nr 2,

pp. 491-497 (USSR)

ABSTRACT:

The authors investigated the reaction between zinc nitrate and NII, in an aqueous medium by isothermal solubility at temperatures of 0° and 25°C. The formation of the basic zinc-nitrate salt at 25°C is terminated at a concentration of 18,77 % NH<sub>3</sub>, 50,14 Zn(NO<sub>3</sub>)<sub>2</sub> and NH<sub>4</sub>(NO<sub>3</sub>). The general formula for the basic zinc-nitrate salt is as follows: Zn(NO<sub>3</sub>)<sub>2</sub>. nZn(OH)<sub>2</sub>. sNH<sub>3</sub>. x H<sub>2</sub>O, where n, s and x have

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different values. The ammoniacal basic salts of zinc

Investigations Concerning the Interaction Between Zinc Salts 78-2-33/43 and Ammonia in an Aqueous Medium

IV. The Solubility of Zinc Nitrate in Aqueous Ammoniacal Solutions

nitrate are highly disperse and by X-ray analysis the same crystal-structure was found for all compounds. In the system  $Zn(NO_3)_2-NH_3-H_2O$  at O C  $Zn(NO_3)_2$ , 4  $NH_3$ , 0,5  $H_2O$  crystallizes and at 25 C  $Zn(NO_3)_2$ , 4  $NH_3$ , 0,25  $H_2O$ . All ammoniacates are resistant to atmospheric influence. The thermographic analyses of tetramine-zinc-nitrate with 0,5 and 0,25 mol  $H_2O$  are equal. In the thermal decomposition three endothermic effects and one exothermic effect occur. At 28-30 C the compounds lose the coarse moisture, at 130 C the crystal water completely escapes and at 200-210 C the melting of anhydrous tetramine-zinc-nitrate occurs. At 205-375 C with an exothermal reaction a spontaneous decomposition of the ammoniacates under formation of 2n0 takes place. There are 8 figures, 3 tables, and 7 references, 3 of which are Slavic.

SUBMITTED:

February 7, 1957

AVAILABLE: Card 2/2 Library of Congress

AUTHORS: -Kirakosyan, A. K., Galustyan, V. D. SOV/78-3-8-33/48

TITLE: The Solubility of Copper Nitrate in Aquecus Ammonia Solutions
(Postvoriment) agentality medical variable and the second of the second o

(Rastvorimost' azotnokisloy medi v vodncammiachnykh rastvorakh)

PERIODICAL: Zhurnal neorganicheskoy khimii, 1958, Vol. 3, Nr. 8, pp. 1925-

1933 (USSR)

ABSTRACT: The solubility of copper nitrate in aqueous ammonia solutions

at 0 and 25°C as well as the solid phases were investigated. The chemical reactions in the copper nitrate, ammonia and water

systems take place in two directions: 1) Crystallization of

basic salts. 2) Crystallization of ammoniates.
The ammonia containing basic salts of corper aget

The ammonia containing basic salts of copper acetate have a variable composition and the same crystallegraphic structure. The basic copper salts may be expressed by the following general formula:  $Cu(NO_3)_2$  •  $nCu(OH)_2$  •  $sNH_3$  •  $xH_2O_3$  s, n, x have dif-

ferent values. The ammonia containing basic copper nitrate salts are fine crystalline deposits; they were investigated by thermal, crystal-optical and x-ray analyses. The thermograms taken display two endothermal effects and one exothermal effect. The endothermal effects occur at 115°C and 185°C and point to a

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SOV/78-3-8-33/48 The Solubility of Copper Nitrate in Aqueous Ammonia Solutions

> dehydration of the basic salts. At 265°C the bonds are decomposed under the formation of copper oxide.

In the system  $Cu(NO_3)_2$ -NH<sub>3</sub>-H<sub>2</sub>O also the ammoniates were crystal-

lized at 0 and 25cc. Monoequotetramine copper nitrate

(Cu(NO3)2.4NH3.H2O) was orystallized. This took place without

difficulties; the compound decomposed under the separation of water and ammonia forming  $Cu(NO_3)_2$  3,5  $NH_3$ . Also the following

compounds were isolated:  $Cu(NO_3)_2 \cdot 4NH_3$ ,  $Cu(NO_3)_2 \cdot 2NH_3$ .

There are 8 figures, 4 tables, and 9 references, 2 of which

are Soviet.

ASSOCIATION: Institut obshchey i neorganicheskoy khimii im. N. S. Kurnakova,

Akademii nauk SSSR (Institute of General and Inorganic Chemistry

imeni N. S. Kurnakov, AS USSR)

SUBMITTED:

June 10, 1957

Card 2/2

URAZOV, G.G. [deceased]; KIRAKOSYAN, A.K.; GAISTYAN, V.D.

Interaction between copper sulfate and ammonia in water. Isv. AN Arm. SSR. Khim. nauki 11 no.4:249-262 '58. (MIRA 11:11)

1. Institut obshchey i neorganicheskiy khimii im N.S. Kurnakova AN SSSR.

(Copper sulfate) (Ammonia)

%(4) AUTHORS:	SOV/76-4-4-25/44 Kirakosyan, A. K., Tananayev, I. V.
TITLE:	Investigation of the Complet Formation of Zirbenium in Schullen by the Ton Execute Method (Izucheniye komplekscobrarovaniya tarrauniya , xastvore s ispol'zovaniyem ionnogo obmena)
FURIDDICAL:	2nurnal meorganicheskoy khimii, 1959 Vol 4, Hr 1, pp 852-856 (USSR)
ABSTRACT:	The authors investigated the complex formation of zircchium sulphate with oxalic, sulphuric and citric acid by means of the cathon exchangers KU-1 and KU-2 under dynamic conditions.
	The complex ion $\left[ Zr(c_2^0)_4 \right]^{4-}$ was found in the system
	2:(SO4)2-H2O2O4-H2O. The compound (2r2O3)O2O4 is fumber at a
	ratio of the components of $H_2C_2O_4$ : $Zr(SO_4)_2 = 0.5$ . The a4-
	sorption of zirconium on both cation exchangers in the NH4",
	No. and H-form in dependence of the exalic acid concent is shown in figures 2 and 3. In the system $Zr(SO_4)_2$ -H_3Cit-E_0
Car: 1/3	zirechium is completely described at the ratio H_Cib:Z=(SC4)2-5

SOV/78-4-4-23/44

Investigation of the Complex Formation of Zirconium in Solution by the Ion Exchange Method

The complex  $\left[\operatorname{Zr}_2(\operatorname{Cit})_7\right]^{X-}$  is produced. At a ratio of the components of  $\operatorname{H}_3(\operatorname{Cit}:\operatorname{Zr}(\operatorname{SO}_4)_2\leqslant 0.34-0.36$  the adsorption of zirconium on the cation exchangers decreases rapidly with increasing ratio of the components. Dizirconyl citrate is probably formed herein. Figure 4 shows the adsorption of zirconium on the cation exchangers in the  $\operatorname{H}^4-$ ,  $\operatorname{Na}^7$ , and  $\operatorname{NH}_4^7-$ form in dependence of the concentration of citric acid. In the system  $\operatorname{Zr}(\operatorname{SO}_4)_2-\operatorname{H}_2\operatorname{SO}_4-\operatorname{H}_2\operatorname{O}$  the zirconyl ion is completely desorbed at a ratio of  $\operatorname{H}_2\operatorname{SO}_4$ :  $\operatorname{Zr}(\operatorname{SO}_4)_2\approx 75$  (concentration of sulphuric acid: 1.5 mols/1). It results from the investigations that direct determination of the composition of the complex ions by the method of ion exchange is only possible with compounds which are in weakly dissociated state present in the solution. There are 5 figures and 7 references, 4 of which are Soviet.

Card 2/3

Investigation of the Complex Formation of Zirconium in Solution of the SOV/78-4-4-23/44 Ion Exchange Method

ASSOCIATION:

Institut obshchey i neorganicheskoy khimii im. N. S. Kurnakova Akademii nauk SSSR (Institute of General and Increaric

Chemistry imeni N. S. Kurnakov of the Academy of Sciences,

USSR)

SUBMITTED:

Jaruary 11, 1958

Card 3/3

Solubility of cadmium bromide in aqueous armonia solutions. Zhur.neorg.khim. 5 no.1:214-218 Ja '60.

(MIRA 13:5)

1. Institut obshchey i neorganicheskoy khimii im. N.S. Kurnakova Ali SSSR.

(Cadmium bromide)

Interaction between sinc sulfate and ammonia in water. Zhur. neorg. khim. 5 no.4:953-959 Ap '60. (MIRA 13:7)

1. Institut obshchey i neorganicheskoy khimii im. H.S. Kurnakova Akademii nauk SSSR.

(Zinc sulfate) (Ammonia)

Solubility of cadmium iodide in aqueous ammonia solutions. Zhurneorg. khim. 5 no.8:1806-1812 Ag \*60. (MIRA 13:9)

1. Institut obshchey i neorganicheskoy khimii im.H.S. Kurnakova Akademii nauk SSSR.

(Cadmium iodide)

S/078/60/005/009/035/040/XX B017/B058

AUTHORS:

Kirakosyan, A. K., Stogova, A. V.

1

TITLE:

Study of the Precipitation Reaction of the Basic Copper

Sulfates With Ammonia

PERIODICAL: Zhurnal neorganicheskoy khimii, 1960, Vol. 5, No. 9, pp. 2088 - 2094

TEXT: The interaction of copper sulfate with ammonia in aqueous phase as well as the composition of the solid phases appearing thereby at 25°C were studied. Five test series were conducted for studying the precipitation reaction of basic copper sulfates. The initial mixtures contained 1.05, 0.5, 0.26, 0.1, and 0.05 mol/l copper sulfate. The ratio of the components NH<sub>3</sub> · CuSO<sub>4</sub> varied from 0.1 to 4.0. The interactions in the system CuSO<sub>4</sub> - NH<sub>3</sub> - H<sub>2</sub>0 at varying initial concentrations of the copper sulfate solutions are tabulated and shown in Figs. 1-5. It follows from the results that the precipitation of basic copper sulfates proceeds in two stages. In the first stage, ammonia acts as basic precipitant and the Card 1/3

Study of the Precipitation Reaction of the S/078/60/005/009/035/040/XX Basic Copper Sulfates With Ammonia B017/B058

reaction product is basic copper sulfate in the second stage, a dissolution of the basic copper sulfate sets in on the addition of an NH<sub>3</sub> excess. The dependence of the pH value (measured by means of the Am-5 (LP-5) tube potentiometer) of the solutions in an equilibrium state on the ratio NH<sub>3</sub>: CuSO<sub>4</sub> in the initial mixture is given in Fig. 6. The dependence of the precipitate composition on the ratio NH<sub>3</sub>: CuSO<sub>4</sub> in the initial mixture is graphically represented in Fig. 7. From the diagrams it can be seen that the precipitated basic copper sulfates have no constant composition in the system CuSO<sub>4</sub> - NH<sub>3</sub> - H<sub>2</sub>O. The more concentrated the initial solutions of copper sulfate, the richer in copper sulfate are the precipitates. Copper hydroxide precipitates predominantly from very diluted solutions of copper sulfate (less than 0.05 mol/1) and at a ratio of the compounds NH<sub>3</sub>: CuSO<sub>4</sub> = 1.5. In the range investigated, two types of basic sulfates were found which crystallize: CuSO<sub>4</sub> nCu(OH)<sub>2</sub> xH<sub>2</sub>O and CuSO<sub>4</sub> nCu(OH)<sub>2</sub> xNH<sub>3</sub> xH<sub>2</sub>O. There are 7 figures, 1 table, and 5 references:

Card 2/3

Study of the Precipitation Reaction of the S/078/60/005/009/035/040/XX Basic Copper Sulfates With Ammonia S/078/60/005/009/035/040/XX

4 Soviet and 1 German.

ASSOCIATION: Institut obshchey i neorganicheskoy khimii im. N.S.Kurnakova Akademii nauk SSSR (Institute of General and Inorganic Chemistry imeni N.S.Kurnakov of the Academy of Sciences USSR)

SUBMITTED: June 9, 1959

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Card 3/3

S/078/60/005/009/036/040/XX B017/B058

AUTHORS: Kirakosyan, A. K., Yeliseyev, A. A.

TITLE: The Interaction of Cadmium Sulfate With Ammonia in the

Aqueous Medium

PERIODICAL: Zhurnal neorganicheskoy khimii, 1960, Vol. 5, No. 9,

pp. 2095 - 2101

TEXT: The properties of basic cadmium sulfates, especially those containing ammonia, were studied by thermal- and X-ray phase analysis. The composition of the basic cadmium sulfates studied is given in Table 1 and the thermograms of these compounds are graphically illustrated in Fig. 1. Two types of basic cadmium sulfates were isolated:  $CdSO_4$  in  $Cd(OH)_2$  in  $H_2O$  and  $CdSO_4$  in  $Cd(OH)_2$  is  $NH_3$  in  $H_2O$ . With a change of the basicity, water- and

ammonia content in the composition of these compounds, a change of their thermal stability and the parameters of the crystal lattices also occurs. The basic cadmium sulfates change their color from white to brown through thermal treatment at temperatures above 150°C. The thermal decomposition

Card 1/2

The Interaction of Cadmium Sulfate With Ammonia in the Aqueous Medium

S/078/60/005/009/036/040/XX B017/B058

of ammonia-containing basic cadmium sulfates differs from that of ammonia-free basic cadmium sulfates. The ammonia-containing basic cadmium sulfates lose the entire water at 400° to 450°C. The X-ray phase analyses proved that all basic cadmium sulfates consist of one phase. The rosntgenograms of the basic cadmium sulfates are shown in Fig. 2 and those of the ammonia-containing basic cadmium sulfates in Fig. 3. The results of the X-ray phase analysis confirm the results of thermal studies. There are 3 figures, 1 table, and 6 references: 5 Soviet and 1 Swiss.

SUBMITTED: June 4, 1959

Card 2/2

KIRAKOSYAN, A.K.

Reaction of precipitation of basic zinc chlorides by ammonia. Zhur. neorg. khim. 6 nc.7:1718-1723 J1 '61. (MIRA 14:7) (Zinc chloride) (Precipitation(Chemistry))

KIRAKOSYAN, A.K.

Reaction of cupric chloride with ammonia. Zhur.neorg.khim. 6

MB:8:1801-1807 Ag '61. (MIRA 14:8)

(Copper chloride) (Ammonia)

S/078/61/006/008/004/018 B121/B203

AUTHORS: Kirakosyan, A. K., Tananayev, I. V.

TITLE: Study of the reaction of zirconium oxychloride with sulfuric acid and with sulfates of ammonium, sodium, iron, and

aluminu

PERIODICAL: Zhurnal neorganicheskoy khimii, v. 6, no. 8, 1961, 1808-1812

TEXT: The authors studied the reactions of  $2rOCl_2$  with  $H_2SO_4$  and with sulfates of ammonium, sodium, iron, and aluminum, as well as the formation of basic zirconium salts by the following methods: Sulfates of the elements mentioned were added in rising amounts to zirconium oxychloride solutions of different concentrations. Studies were made at room temperature. The equilibrium between liquid and solid phase was established within 5 - 30 days as dependent on the content of oxychloride in the initial mixture. The reaction of zirconium oxychloride with sulfuric acid, and with sulfates of socium, ammonium, iron, and aluminum in aqueous phase is supposed to proceed in three stages: In the first stage, soluble basic zirconium sulfates are obviously formed which only precipitate at pH = 1.

Card 1/3

Study of the reaction...

S/C78/61/CO6/CO8/CO4/U18 B121/B203

ASSOCIATION: Institut obshchey i neorganicheskoy khimii im. N. S. Kurnakova Akademii nauk SSSR (Institute of General and Inorganic Chemistry imeni N. S. Kurnakov of the Academy of Sciences USSR)

SUBMITTED: June 15, 1960

Card 3/3

## EXECUTION OF Basic cadmium chlorides by ammonia. Precipitation of basic cadmium chlorides by ammonia. Zhur.neorg.khim. 7 no.11:2557-2562 N '62. (MIRA 15:12) (Cadmium chloride) (Ammonia)

Hasic zinc sulfates. Zhur.neorg.khim. 8 no.1:119-129 Ja 163.

(HIRA 16<sup>15</sup>)

(Zinc sulfates)

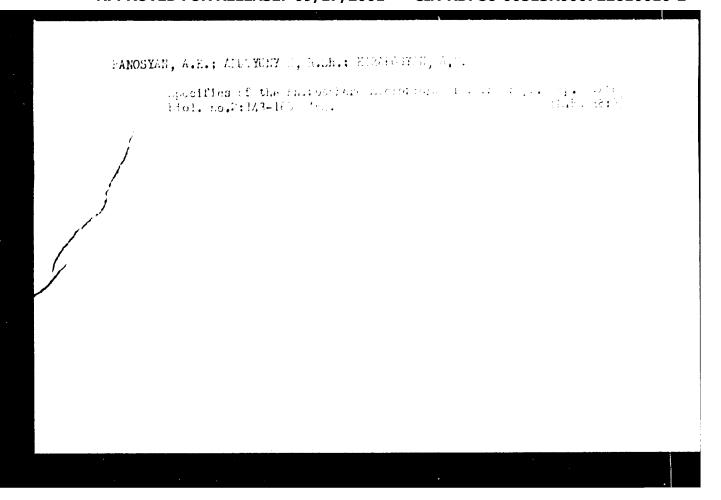
# Frecipitation of basic cadmium bromides by ammonia. Zhur.neorg.khim. 8 no.31622-628 Mr 163. (MRA 16:4) (Cadmium bromide) (Ammonia)

## KIRAKOSYAN, A.K.

Precipitation reactions of copper subbromides by ammonia. Zhur.neorg.khim. 8 no.4:905-910 Ap '63. (MIRA 16:3)
(Copper bromides) (Ammonia) (Precipitation (Chemistry))

KIRAKOSYAN, A.K.; RADOSTINA, L.B.

Reaction of the precipitation of basic zinc bromides by ammonia. Zhur. neorg. khim. 10 no.1:160-165 Ja \*65. (MIRA 18:11) 1. Submitted April 12, 1963.

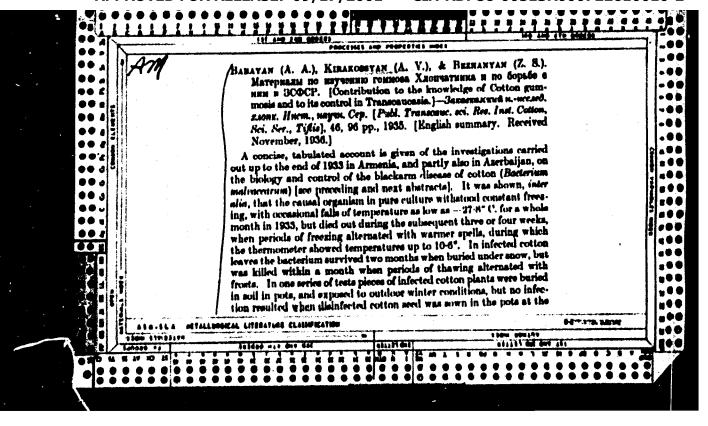


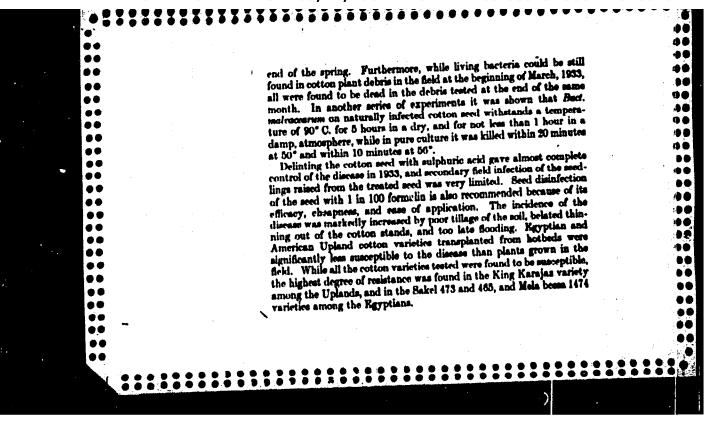
## KIRAKOSYAN, A.S. Impossibility of economic planning under capitalism [in Armenian with summary in Russian]. Nauch.trudy Erev.un. 56:43-69 '56. (MIRA 10:7)

1. Kafedra planirovaniya narodnogo khozyaystva. (Economic policy)

## KIRAKOSYAN, Amushavan Saribekovich

[Planning of the supply of materials and equipment] [Planivanie material\*no-tekhnicheskogo snebzheniia. Erevan, Izd-vo "Mitk"] 1965. 182 p. [In Armenian] (MIRA 18:8)





## PETROSYAN, A.P.; KIRAKOSYAN, A.V.

Specificity of Azotobacter for various agricultural crops. Mikrobiol.sbor. no.4:25-42.149.
(AZOTOBACTER)

(AZOTOBACTER)

KIRAKOSYAH, A.V.; PETROSYAH, A.P.; AZARYAH, E.Kh.

Effect of the bacteria of activators on the effectiveness of Azotobacter. Mikrobiol.abor. no.4:43-65 '49. (MLRA 9:8) (RHIZOSPHERE MICROBIOLOGY) (AZOTOBACTER)

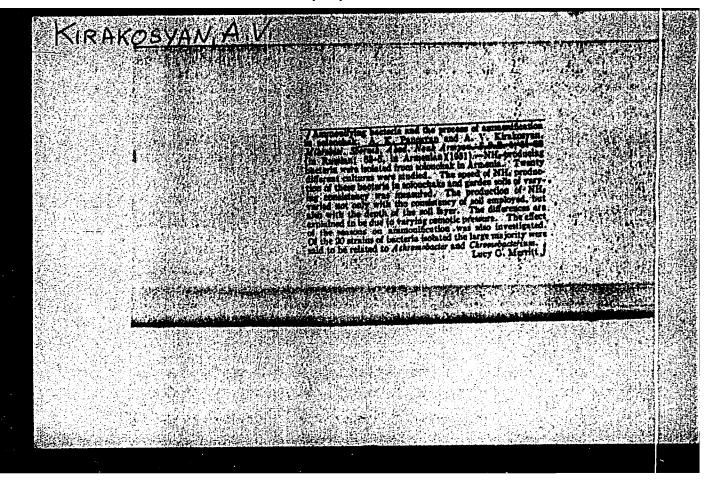
## KIRAKOSYAN, A.V.

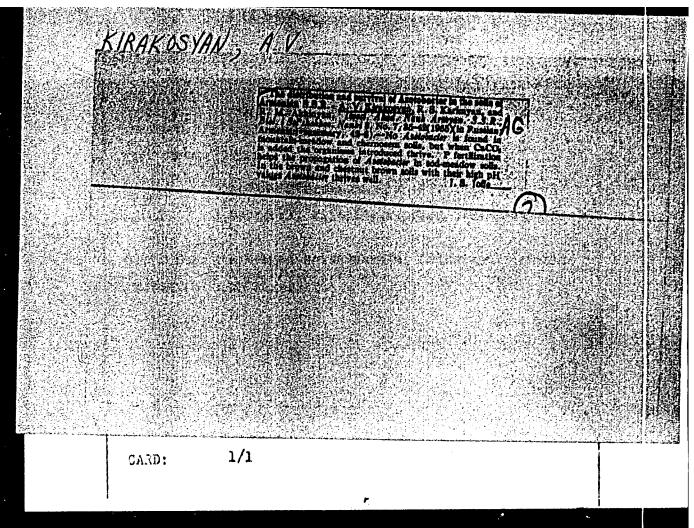
Development of micro-organisms in the rhizosphere of some agricultural crops. Mikrobiol.shor. no.4:67-83 '49. (MIRA 9:8) (RHIZOSPHERE MICROBIOLOGY)

KIRAKOSYAN, A. V.

Kirakosyan, A. V. and Khachatrian, G. A. "Virus Diseases of Potatoes in Armenia," <u>Isvestiia Akademii Nauk Armianskoi SSR</u>, vol. 3, 1950, pp. 33-334. 20 Erk

SO: SIRA S. 19-53, 15DEC 1953





. KIRAKUSYAN A.V

USSR / Microbiology. Antibiosis and Symbiosis. Antibiotics. Antibiosis.

F.

Abs Jour : Ref Zhur - Biologiya, No 6, 1959, No. 24026

Author : Kirakosyan, A. V. Karimyan, R. S.

Inst : Not given
Title : Intraspecific and Interspecific Interrelations

of Azotobacter

Orig Pub : Mikrobiol. sb. All ArmSSR, 1958, vyp 9, 3-22

Abstract: The intrespecific and interspecific interrelations were studied in 280 cultures of
azotobacter, isolated from various types of
soil of the Armenian SSR (190 cultures of
Azotobacter chrococcum, 64 of A. nigricans,
21 of A. agile and 3 of A. vinelandii).

Antagonistic Interrelations were discovered

not only between the various types of azotobacter

Card 1/3

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APPROYED FOR RELEASE: 09/17/2001 and SJA5ROR86-00513R000722610010-2"
Antibiotics. Antibiosis.

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but also between the various strains of one and the same type. 32% of tested cultures manifested intraspecific antagonistic action. The largest percentage of intraspecific antagonists was discovered among the representatives of species of Az. chroococcum. The cultures of azotobacter with strong antagonistic action are usually antagonists with respect to the greatest number of cultures intraspecifically, as well as among other types of azotobacter, and are themselves, as a rule, rarely subject to antagonistic action of other cultures of azotobacter. No correlation was discovered between the type of soil and the presence of antagonistic properties in

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